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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO	
10/827,106 04/19/2004		Gopi M. Venkatesh	451194-107	1448	
27805 75	1111515000		EXAMINER		
THOMPSON HINE L.L.P. P.O. BOX 8801			SAMALA, JAGADISHWAR RAO		
DAYTON, OH 45401-8801			ART UNIT	PAPER NUMBER	
			1618	:	
			DATE MAILED: 11/13/2006		

Please find below and/or attached an Office communication concerning this application or proceeding.

		Applie	cation No.	Applicant(s)					
Office Action Summary		10/82	7,106	VENKATESH ET AL.					
		Exam	iner	Art Unit					
		Jagad	ishwar R. Samala	1618					
Period fo	The MAILING DATE of this commun r Reply	ication appears or	the cover sheet with the	correspondence ac	ldress				
WHIC - Exter - after - If NO - Failu Any r	ORTENED STATUTORY PERIOD F CHEVER IS LONGER, FROM THE M Isions of time may be available under the provisions SIX (6) MONTHS from the mailing date of this come period for reply is specified above, the maximum stars to reply within the set or extended period for reply reply received by the Office later than three months and patent term adjustment. See 37 CFR 1.704(b).	AALLING DATE OF of 37 CFR 1.136(a). In r nunication. atutory period will apply a v will, by statute, cause the	THIS COMMUNICATIO no event, however, may a reply be ti nd will expire SIX (6) MONTHS from a application to become ABANDON	N. mely filed n the mailing date of this c ED (35 U.S.C. § 133).					
Status									
1)	Responsive to communication(s) file	ed on							
•		2b)⊠ This action	is non-final.						
	<u> </u>								
٠,٠	closed in accordance with the practice under Ex parte Quayle, 1935 C.D. 11, 453 O.G. 213.								
Dispositi	on of Claims	·							
4)⊠ Claim(s) <u>1-24</u> is/are pending in the application.									
-	4a) Of the above claim(s) is/are withdrawn from consideration.								
	5) Claim(s) is/are allowed.								
·	☐ Claim(s) 1-24 is/are rejected.								
8)□	Claim(s) are subject to restrict	ction and/or election	on requirement.						
Applicati	on Papers								
_	·	e Examiner							
9) The specification is objected to by the Examiner. 10) The drawing(s) filed on is/are: a) accepted or b) objected to by the Examiner.									
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).									
	Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).								
11)	11) The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.								
Priority u	ınder 35 U.S.C. § 119								
12)	Acknowledgment is made of a claim	for foreign priority	under 35 U.S.C. § 119(a	a)-(d) or (f).					
	☐ All b)☐ Some * c)☐ None of:	5 1 3	•						
	1. ☐ Certified copies of the priority documents have been received.								
	2. Certified copies of the priority documents have been received in Application No								
	3. Copies of the certified copies of the priority documents have been received in this National Stage								
•	application from the Internation	nal Bureau (PCT	Rule 17.2(a)).						
* S	See the attached detailed Office action	on for a list of the o	ertified copies not receiv	ed.					
Attachmen									
1) 🔯 Notic 2) 🔲 Notic	e of References Cited (PTO-892) e of Draftsperson's Patent Drawing Review (F	OTO 048\	4) Interview Summar Paper No(s)/Mail D						
3) 🔯 Inform	nation Disclosure Statement(s) (PTO/SB/08) r No(s)/Mail Date 11/19/04; 12/19/05		5) Notice of Informal 6) Other:						

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DETAILED ACTION

Application Status

1. Applicant's response to the Office Action was acknowledged April 19, 2004

Claim Disposition

2. Claims 1-24 is/ are pending and are under examination.

Information Disclosure Statement

4. The Information Disclosure Statement filed on November 19, 2004; December 19, 2005 and May 08, 2006 has been received and entered the references cited on the PTO-1449 Form have been considered by the examiner and a copy is attached to the instant Office Action.

Claim Rejections - 35 USC § 102

1. The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless -

- (b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.
- 2. Claims 1-24 are rejected under 35 U.S.C. 102(b) as being anticipated by Ohta, Motohiro et al., (EP 0 914818 A1 here after '818).

Claims 1-24 are drawn to a tablet composition and method of manufacturing tablet that disintegrates in the oral cavity comprising a compressed blend of a rapidly

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dispersing microgranules prepared by granulating a sugar alcohol or saccharide having an average particle size less than 30 microns, a disintegrant and a taste masked microcapsule containing at least one drug.

The patent '818 discloses a production method of a tablet characterized by compressing powdered mixture comprising sugar alcohol or saccharide each having an average particle diameter of not more than 30 microns ground by means of a hammer mill or a jet mill or the like, an active ingredient, and a disintegrant under the presence of an easily volatile disintegrating adjuvant and thereafter the disinitegrant adjuvant is volatilized. (see 0004) The disintegrant mainly used such as crospovindone, crosscarmellose sodium, low substituted hydroxypropylcellulose or the like which is widely used for drugs and food (see 0016). Also sugar alcohol used were D-mannitol, sorbitol, and saccharide used were lactose and glucose or like which is widely used for drugs and foods. The amount of sugar alcohol or saccharide is preferably about 60-95 % by weight of tablet (see 006 and 0019). The amount of taste masked active ingredient is 0.01-30 %, and the amount of disintegrant present is preferably about 1-30mg per dosage, and more preferably 1-10 % per one tablet (see 0021). The patent '818 also discloses the use of binders such as hydroxypropylcellulose, polyvinylpyrrolidone, hydroxymethlylcellulose, partially saponificated polyvinyl alcohol methylcellulose, pullulan or the like, sour agent as citric acid, malic acid, adipic acid ascorbic acid or like, sweetening agent is Aspartame ™, saccharin, glycyrrhizic acid or the like, flavoring agent is lemon, orange, pine, mint, menthol or the like, colorant is vellow iron sesquioxide, red iron sesquioxide, tar color or the like optionally as additional

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pharmaceutical ingredients (see 0023). The patent further discloses the following drugs such as hyprotic, anxiolytic, antiepileptic, psychoneurosis agents, drug for central nervous system, peripheral nervous system, autonomic nervous system, skeletal muscle relaxant, antiarrhythmic agent, angiotonic, vasodilator, hyperlipemia, antidiarrheal drug, drug for peptic ulcer such as azulene, L-glutamine, aceglutamide aluminium, cetraxate hydrochloride, cimetidine and so on (see 0007 –0015). The teaching of the patent '818 possesses all the steps required as recited in claims. Thus, all the claims are anticipated.

Claim Rejections - 35 USC § 103

- 3. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:
 - (a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.
- 4. Claims 1-24 are rejected under 35 U.S.C. 103(a) as being unpatentable over Percel et al. (US 6,451345 here after '345) in view of Masaki et al. (US 5,466,464 here after '464).

Percel discloses a taste masked antibiotic tablet comprising a primary ethylcellulose coating of Linezolid needle-like crystals with a median particle size of 20 microns by solvent coacervation, an optionally blended with other pharmaceutically acceptable excipients such as binders like hydroxypropylcellulose, microcrystalline cellulose, disintegrants include cornstarch, lactose, mannitol, sucrose, crospovidone,

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flavors, sweeteners, suspending agents and/or preservatives and filled into unit dose containers or compressed into fast-disintegrating/effervescent/chewable tablets.(see abstract, column 1, lines 17-24 and column 5, example 7).

Percel fails to disclose H_2 antagonists such as ranitidine, cimetidine, and famotidine and so on as a pharmaceutically active ingredient included therein. However, the use of H_2 –receptor antagonist useful for inhibiting gastric secretion and treating ulcers is well known in the art as shown by Masaki.

Masaki discloses a process for the production of an intrabuccally disintegrating solid tablet preparation comprising an active ingredient (i.e. H₂-receptor antagonists such as famotidine, cimetidine, ranitidine, and a sugar comprising lactose and/or mannitol (see column 3, lines 7-20).

It would have been obvious to one of ordinary skill in the art to modify the Parcel chewing tablets to include an active medicament, H₂-receptors as an additional pharmaceutical active medicamnet because Masaki teaches that addition of H₂-receptor antagonists as a active medicament greatly reduces gastric acid secretion systemically and healing ulcers. One of ordinary skill in the art would have been motivated to include the H₂-receptor antagonists as an additional pharmaceutical active medicament disclosed by Parcel because H₂-receptors antagonists taught by Masaki, while having similar physiological functions, provides an additional treatment for upper abdominal pain/discomfort and heartburn, indigestion, sour stomach, and other gastrointestinal disorders including gastro-oesophageal reflux.

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Conclusion

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1. No claims are allowed at this time.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Jagadishwar R. Samala whose telephone number is (571)272-9927. The examiner can normally be reached on 8.30 A.M to 5.00 P.M.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Michael G. Hartley can be reached on (571)272-0616. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

VICKIE KIM
PRIMARY EXAMINER

Jagadishwar R Samala Examiner Art Unit 1618